**AMENDMENTS TO THE CLAIMS** 

1. (Currently amended) A method for identifying an agent capable of modulating

expression of CYP2S1 by a cell, comprising the steps of:

a) providing a cell or cells capable of expressing CYP2S1 comprising a reporter gene or

CYP2S1 gene under control of a regulatory sequence shown in Figure 7 comprising at least an

XRE-like sequence, an AP-1-like sequence or a RARE-like sequence;

b) contacting a test agent with said cell(s);

c) incubating said cell(s) under conditions which are conducive to enable expression of

said CYP2S1 gene or reporter gene when in the absence of the test agent; and

d) detecting whether or not the test agent modulates modulation of expression of said

CYP2S1 gene or said reporter gene.

2. (Cancelled)

3. (Currently amended) The method according to claim [[2]] 1 wherein the reporter

nucleic acid is capable of encoding gene encodes glutathione S-transferase, an antibiotic, a

chromogenic substrate, such as  $\beta$ -galactocidase, luciferase, a fluorescent protein, such as green

fluorescent protein, or chloramphenicol acetyl transferase.

4. (Currently amended) The method according to any preceding claim 1 wherein said

cell(s) is/are a skin cell(s).

5. (Cancelled)

6. (Currently amended) The method according to any of claims 1 to 3 claim 1 wherein

said cell(s) is a mammalian, mammalian or bacterial, yeast or insect cell which has been genetically

engineered so as to be capable of expressing said CYP2S1 gene or said reporter nucleic acid gene.

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7. (Currently amended) The method according to claim [[6]] 4, when dependent on

claim 1 wherein the cell(s) has/have been genetically engineered so as to comprise a nucleic acid

capable of encoding CYP2S1 and a sequence upstream thereof as shown in Figure 7 capable of

controlling transcription and/or translation of said nucleic acid.

8-14. (Cancelled)

15. (Currently amended) The method according to any one of claims 1 and 4 to 8 claim

39 wherein detection of any modulation in the expression of CYP2S1 is carried out using an

antibody specifically reactive to CYP2S1.

16. (Currently amended) The method according to any one of claims 1 and 4 to 8 claim

39 wherein detection of any modulation in the expression of CYP2S1 mRNA is carried out using

quantitative real time PCR analysis.

17. (Cancelled)

18. (Currently amended) A recombinant expression vector comprising a nucleic acid

capable of encoding CYP2S1 or a reporter protein under transcriptional and/or translational control

of the an isolated nucleic acid molecule according to claim 17 comprising the regulatory sequence

shown in Figure 7.

19. (Cancelled)

20. (Currently amended) A host cell comprising the recombinant vector according to

either of claims 18 or 19 claim 18.

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21. (Currently Amended) The host cell according to claim 20 wherein the cell is a

mammalian[[,]] or bacterial, yeast or insect cell which has been genetically engineered so as to be

capable of expressing CYP2S1 or said reporter nucleic acid.

22. (Cancelled)

23. (Original) A method of making CYP2S1 comprising culturing the host cell according

to claim 20 under conditions such that CYP2S1 is expressed; and recovering CYP2S1.

24. (Cancelled)

25. (Currently amended) A pharmaceutical composition comprising isolated CYP2S1

according to claim 24 in combination with a pharmaceutically acceptable carrier therefore.

26-31. (Cancelled)

32. (Currently amended) A method of preventing, treating or ameliorating in a subject a

skin condition in a subject related to increased or decreased CYP2S1 expression in skin, which

comprises administering to a mammalian the subject CYP2S1, a vector capable of expressing

CYP2S1, or an agent capable of modulating expression of CYP2S1 in skin tissue.

33. (Currently amended) A method of diagnosing a skin condition associated with

increased or decreased expression of CYP2S1, or a predisposition to such a skin condition

comprising detecting a level of CYP2S1 in a test skin sample according to the method of claim 39

and comparing this said level against a normal control, wherein such that an increase or decrease in

the CYP2S1 expression level in the test skin sample as compared to the normal control is indicative

of [[a]] said skin condition or said predisposition to a skin condition.

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34. (Currently amended) A method of diagnosing a skin condition associated with

increased or decreased expression of CYP2S1 or a predisposition to such a skin condition associated

with increased or decreased expression of CYP2S1, comprising detecting a polymorphism in a

CYP2S1 gene or upstream sequence thereof, which effects affects expression of CYP2S1, wherein

detection of a polymorphism is indicative of [[a]] said skin disorder associated with increased or

decreased CYP2S1 expression, or said predisposition thereto.

35. (Currently amended) A method of detecting effectiveness of a skin treatment to be

administered to a patient suffering from a skin condition, comprising the steps of:

a) obtaining a <u>first</u> sample of diseased skin <u>from the patient</u> and detecting <u>according to</u>

the method of claim 39 a level of CYP2S1 expression in the first sample of diseased skin prior to

administration of the skin treatment;

b) administering said skin treatment to the patient; and

c) after a period of time, obtaining a further second sample of diseased skin from the

patient and detecting according to the method of claim 39 whether or not there has been an increase

or decrease in the level of CYP2S1 compared to the first sample expression.

36. (Currently amended) A method of detecting whether or not a subject is likely to

respond to a skin treatment with a chemical which is metabolisable by CYP2S1, comprising the

steps of:

a) obtaining samples a first sample of diseased skin and a second sample of non-

diseased skin from a subject; and

b) detecting according to the method of claim 39 a level of CYP2S1 expression in the

diseased and non-diseased first and second samples

wherein an increase in expression of the CYP2S1 level in diseased skin the first sample compared to

the second sample is indicative of a subject who may respond favourably to a chemical which is

metabolisable by CYP2S1 said skin treatment.

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37. (Currently amended) A method of identifying possible a new skin treatment drug

candidates candidate comprising contacting the drug candidate with CYP2S1 and observing for

detecting metabolites of said drug candidate.

38. (Currently amended) A method of improving effectiveness of a skin treatment being

administered to a subject comprising the steps of

a) detecting according to the method of claim 39 a level of CYP2S1 in the skin of said

subject; and

b) either increasing or decreasing expression the level of CYP2S1 in diseased the skin to-be

treated of said subject receiving said skin treatment.

39. (New) A method of detecting a level of CYP2S1 in a skin cell, comprising

determining the level of expression of CYP2S1 in said skin cell.